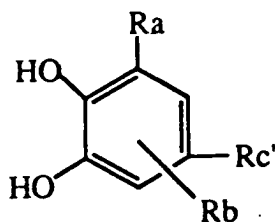


CM 40. A compound of the formula



Ib

CF5 wherein Ra is nitro or cyano, Rb is hydrogen or halogen, Rc' is the group CO-R<sup>11</sup> wherein R<sup>11</sup> is a phenyl group optionally mono- or disubstituted by halogen, cyano, hydroxy or lower alkyl, or an ester or ether derivative thereof which is hydrolyzable under physiological conditions or a pharmaceutically acceptable salt thereof.

2  
Claim 2, line 1: please change "Claim 1" to  
--- Claim 40 ---.

Claim 8, line 1: please change "Claim 1" to  
--- Claim 40 ---.

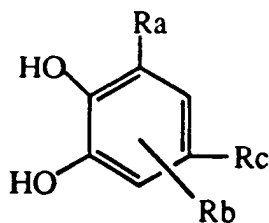
P  
Claim 9, line 1: please change "Claim 1" to  
--- Claim 40 ---.

Please cancel Claim 10.

2  
K  
Please cancel claim 11 and replace with new Claim 41.

8 ~~41~~. A pharmaceutical composition comprising a compound of the formula

T1130X



02  
PS wherein Ra is nitro or cyano, Rb is hydrogen or halogen, Rc is CO-R'<sup>13</sup> wherein R' is a phenyl group optionally mono- or disubstituted by halogen, cyano, hydroxy or lower alkyl or an ester or ether derivative thereof which is hydrolyzable under physiological conditions or a pharmaceutically acceptable salt thereof, and a therapeutically inert carrier material.

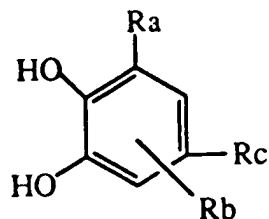
2  
P  
K  
Claim 12, line 2: please change "11" to --- 41 ---.

Claim 13, line 2: please change "11" to --- 41 ---.

Please cancel Claim 14.

Please cancel Claim 15 and replace with new Claim 42.

42. A pharmaceutical composition comprising L-dopa,  
peripheral decarboxylase inhibitor, a compound of the formula



PS wherein Ra is nitro or cyano, Rb is hydrogen or halogen, Rc is the group CO-R<sup>1</sup> wherein R<sup>1</sup> is a phenyl group optionally mono- or disubstituted by halogen, cyano, hydroxy or lower alkyl or an ester or ether derivative thereof which is hydrolyzable under physiological conditions or a pharmaceutically acceptable salt thereof and a therapeutically inert carrier material.

Claim 16, line 2: please change "15" to --- 42 ---.

Claim 17, line 15: please change "15" to --- 42 ---.

Please cancel Claims 18-30.

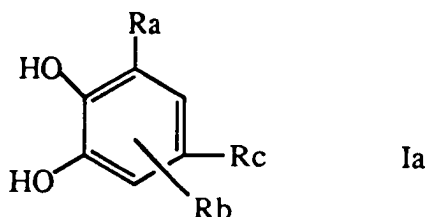
Claim 31, line 1: please change "Claim 1" to  
--- Claim 40 ---.

Claim 32, line 1: please change "Claim 11" to  
--- Claim 41 ---.

Claim 33, line 1: please change "Claim 15" to  
--- Claim 42 ---.

Please cancel Claim 34 and replace with new Claim 43.

<sup>17</sup>  
~~43.~~ A pharmaceutical composition for treating depression  
comprising an effective amount of a compound of the formula

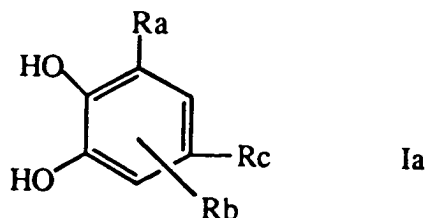


CY  
wherein Ra is nitro or cyano, Rb is hydrogen or halogen, Rc is  
CO-R<sup>1</sup> wherein R<sup>1</sup> is a phenyl group optionally mono- or  
disubstituted by halogen, cyano, hydroxy or lower alkyl or an  
ester or ether derivative thereof which is hydrolyzable under  
physiological conditions or a pharmaceutically acceptable salt  
thereof, and a therapeutically inert carrier material.

Claim 35, line 1: please change "Claim 34" to  
--- Claim 43 ---.

Please cancel Claim 36 and replace with new Claim 44.

<sup>19</sup>  
44. A pharmaceutical composition for treating Parkinson's disease comprising L-dopa, a peripheral decarboxylase inhibitor, a compound of the formula

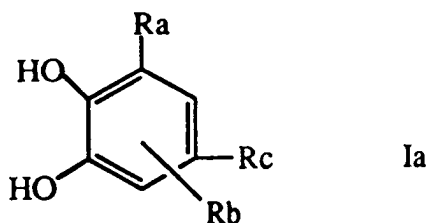


wherein Ra is nitro or cyano, Rb is hydrogen or halogen, Rc is CO-R<sup>1</sup> wherein R<sup>1</sup> is a phenyl group optionally mono- or disubstituted by halogen, cyano, hydroxy or lower alkyl or an ester or ether derivative thereof which is hydrolyzable under physiological conditions or a pharmaceutically acceptable salt thereof, and a therapeutically inert carrier material.

Claim 37, line 1: please change "Claim 36" to  
--- Claim 44 ---.

Please cancel Claim 38 and replace with new Claim 45.

<sup>2/</sup>  
45. A pharmaceutical composition for inhibiting catechol-O-methyl-transferase, said composition comprising a catechol-O-methyl-transferase inhibiting amount of a compound of the formula



wherein Ra is nitro or cyano, Rb is hydrogen or halogen, Rc is CO-R<sup>1</sup> wherein R<sup>1</sup> is a phenyl group optionally mono- or disubstituted by halogen, cyano, hydroxy or lower alkyl, or an ester or ether derivative thereof which is hydrolyzable under physiological conditions or a pharmaceutically acceptable salt thereof, and a therapeutically inert carrier material.

Claim 39, line 1: please change "Claim 38" to  
--- Claim 45 ---.

REMARKS

Reconsideration of the captioned application in view of the foregoing amendments and the remarks which follow is courteously requested.